Proteins

Product Data Sheet

PROTAC Bcl-xL degrader-2

Cat. No.: HY-139309 Molecular Formula: $C_{68}H_{80}N_8O_{14}S_3$

Molecular Weight: 1329.6

Bcl-2 Family; PROTACs Target: Apoptosis; PROTAC Pathway:

Storage: Powder -20°C 3 years

In solvent

4°C 2 years -80°C 6 months

-20°C 1 month



SOLVENT & SOLUBILITY

In Vitro

DMSO: 100 mg/mL (75.21 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	0.7521 mL	3.7605 mL	7.5211 mL
	5 mM	0.1504 mL	0.7521 mL	1.5042 mL
	10 mM	0.0752 mL	0.3761 mL	0.7521 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

1. Add each solvent one by one: 10% DMSO >> 90% corn oil

Solubility: 10 mg/mL (7.52 mM); Suspended solution; Need ultrasonic

BIOLOGICAL ACTIVITY

Description PROTAC Bcl-xL degrader-2 is a potent Bcl-xL (Bcl-2 family member) degrader based on von Hippel-Lindau ligand, with an IC ₅₀ of 0.6 nM.

Bcl-xL VHL IC₅₀ & Target

0.6 nM (IC₅₀)

PROTAC Bcl-xL degrader-2 (PROTAC 6; 0.1 nM-10 µM; 24 hours) treatment decreases the level of Bcl-xL protein in THP-1 cells In Vitro

PROTAC Bcl-xL degrader-2 (PROTAC 6) inhibits caspase 3/7 activity in MOLT-4 cells with an IC₅₀ of 466 nM^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Western Blot Analysis $^{[1]}$

Cell Line: THP-1 cells

Concentration:	0.1 nM-10 μM
Incubation Time:	24 hours
Result:	Led to monotonic decreases in observed Bcl-xL protein.

REFERENCES

[1]. Chun-Wa Chung, et al. Structural Insights into PROTAC-Mediated Degradation of Bcl-xL. ACS Chem Biol. 2020 Sep 18;15(9):2316-2323.

Caution: Product has not been fully validated for medical applications. For research use only.

Tel: 609-228-6898 Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA

Page 2 of 2 www.MedChemExpress.com